Application No. 10/666,642 Amendment dated February 13, 2006 Reply to Office action of October 11, 2005

Amendments to the Claims:

This listing of claims will replace all prior versions and listings of claims in the application.

Listing of Claims:

Claims 1-20 (canceled)

Claim 21 (previously presented): A transgenic plant transformed with an expression vector comprising a polynucleotide sequence encoding a polypeptide having a conserved domain that has at least 70% sequence identity to the conserved domain of amino acid coordinates 111-164 of SEQ ID NO: 194.

Claim 22 (previously presented): The transgenic plant of claim 21, wherein the expression vector further comprises a constitutive, inducible, or tissue-specific promoter operably linked to the polynucleotide sequence.

Claim 23 (currently amended): Seed A seed produced by the transgenic plant according to claim 21, wherein the seed comprises the polynucleotide sequence of claim 21.

Claim 24 (currently amended): A method for producing a transgenic plant having an altered trait as compared to a wild-type plant of the same species, the method steps comprising:

- (a) providing an expression vector comprising:
 - (i) a polynucleotide sequence encoding a polypeptide comprising a conserved domain that has at least 70% sequence identity to the conserved domain of amino acid coordinates 111-164 of SEQ ID NO: 194; and
 - (ii) at least one regulatory element operably linked to the polynucleotide sequence, wherein said at least one regulatory element controls expression of the polynucleotide sequence in a target plant;
- (b) introducing the expression vector into at least one plant; and
- (c) selecting at least one transgenic plant that has an altered trait as compared to a wild-type plant of the same species;

wherein the altered trait is selected from the group consisting of greater tolerance to cold during germination, greater tolerance to cold during growth, greater tolerance to drought water deprivation, greater tolerance to nitrogen limitation, larger leaves, and greater biomass than the wild-type plant.

Claim 25 (previously presented): The method of claim 24, wherein the polypeptide comprises a conserved

domain that has at least 75% sequence identity to the conserved domain of amino acid coordinates 111-164

of SEQ ID NO: 194.

Claim 26 (previously presented): The method of claim 24, wherein the polypeptide comprises a conserved

domain that has at least 80% sequence identity to the conserved domain of amino acid coordinates 111-164

of SEQ ID NO: 194.

Claim 27 (previously presented): The method of claim 24, wherein the regulatory element is a cauliflower

mosaic virus 35S promoter.

Claim 28 (previously presented): The method of claim 24, wherein the regulatory element is a root-specific,

epidermis-specific, meristem-specific, or leaf-specific promoter.

Claim 29 (previously presented): The method of claim 24, wherein the regulatory element is a drought-

inducible or cold-inducible promoter.

Claim 30 (canceled)

Claim 31 (previously presented): The method of claim 24, wherein the transgenic plant has greater tolerance

to 168 hours without watering than the wild-type plant.

Claim 32 (previously presented): The method of claim 24, wherein the transgenic plant has greater tolerance

than the wild-type plant to six hours of exposure to 8° C during germination or six hours of exposure to 4°

to 8° C during growth.

Claim 33 (previously presented): The method of claim 24, wherein the transgenic plant has greater tolerance

than the wild-type plant to MS media with the nitrogen source reduced to 20 mg/l of NH₄NO₃.

Claim 34 (currently amended): Seed A seed produced by a transgenic plant produced by the method

according to claim 24, wherein the seed comprises the expression vector of claim 24.

Claim 35 (currently amended): A method for increasing the tolerance of a plant to an abiotic stress, the

method steps comprising:

Page 11 of 23

Application No. 10/666,642 Amendment dated February 13, 2006 Reply to Office action of October 11, 2005

(a) providing an expression vector comprising:

(i) a polynucleotide sequence encoding a polypeptide comprising a conserved domain that has at least 70% sequence identity to the conserved domain of amino acid coordinates 111-164 of

SEQ ID NO: 194; and

(ii) at least one regulatory element flanking the polynucleotide sequence, wherein said at least one regulatory element controls expression of the polynucleotide sequence in a target plant;

(b) introducing the expression vector into a plant, thereby producing a transgenic plant; and

(c) selecting a transgenic plant having an altered trait as compared to a wild-type plant of the same

species;

wherein the abiotic stress is selected from the group consisting of cold stress during germination,

cold stress during growth, drought stress water deprivation, and nitrogen limitation.

Claim 36 (previously presented): The method of claim 35, wherein the polypeptide comprises a conserved

domain that has at least 75% sequence identity to the conserved domain of amino acid coordinates 111-164

of SEQ ID NO: 194.

Claim 37 (previously presented): The method of claim 35, wherein the polypeptide comprises a conserved

domain that has at least 80% sequence identity to the conserved domain of amino acid coordinates 111-164

of SEQ ID NO: 194.

Claim 38 (previously presented): The method of claim 35, wherein the transgenic plant has greater tolerance

to 168 hours without watering than the wild-type plant.

Claim 39 (previously presented): The method of claim 35, wherein the transgenic plant has greater tolerance

than the wild-type plant to six hours of exposure to 8° C during germination or six hours of exposure to 4°

to 8° C during growth.

Claim 40 (previously presented): The method of claim 35, wherein the transgenic plant has greater tolerance

than the wild-type plant to MS media with the nitrogen source reduced to 20 mg/l of NH₄NO₃.

Page 12 of 23